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# Modelling of a Biomass Gasification Plant Feeding a Hybrid Solid Oxide Fuel Cell and Micro Gas Turbine System

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## Abstract

A system level modelling study on two combined heat and power (CHP) systems both based on biomass gasification. One system converts the product gas in a solid oxide fuel cell (SOFC) and the other in a combined SOFC and micro gas turbine (MGT) arrangement. An electrochemical model of the SOFC has been developed and calibrated against published data from Topsoe Fuel Cells A/S (TOFC) and Risø National Laboratory. The modelled gasifier is based on an up scaled version of the demonstrated low tar gasifier, Viking, situated at the Technical University of Denmark. The MGT utilizes the unconverted syngas from the SOFC to produce more power as well as pressurizing the SOFC bettering the electrical efficiency compared to operation with the SOFC alone - from  $\eta_{el}=36.4\%$  to  $\eta_{el}=50.3\%$ .

*Keywords:* System modelling, biomass gasification, micro gas turbine, SOFC

## Nomenclature

$a_{ohm}, b_{ohm}$	coefficients for Eq. (24)
$ASR$	area specific resistance
$E$	reversible open circuit voltage
$F$	Faradays constant
$g_f$	Gibbs free energy of formation
$i$	current density
$LHV$	lower heating value
$\dot{n}$	molar flow
$n_e$	transferred electrons per molecule of fuel
$p$	pressure/partial pressure
$P$	power production
$R$	universal gas constant
$T$	temperature
$UF$	fuel utilization factor for SOFC
$V$	potential/overpotential
$y$	molar fraction
$\delta$	SOFC layer thickness
$\eta$	efficiency

Subscripts:

a	anode
c	cathode
con	consumption
e	electrolyte
i	interconnect

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# 1 Introduction

Development of sustainable and efficient production plants of combined heat and power (CHP) tends to gain more attention as climate changes, security of supply and depletion of fossil fuels have become well known issues. The share of biomass in CHP production are expected to increase in the future and decentralized CHP plants are also of interest to avoid costs of biomass transportation. Efficient power producing technologies for small scale productions are typically gas engines, micro gas turbines (MGT) and fuel cells – all requiring gaseous fuel. Gasification can deliver biomass based gaseous fuel so the combination of biomass gasification and efficient syngas conversion are potentially a sustainable and efficient CHP plant.

Solid oxide fuel cells (SOFCs) can electrochemically convert  $H_2$  and CO as well as internally reform  $CH_4$  into more  $H_2$  and CO due to their high operating temperature. This makes SOFCs very fuel flexible and ideal for converting syngas compared to other fuel cell types.

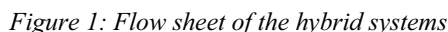
The performance and system design of integrated biomass gasifier and SOFC systems in the 100-1000kW<sub>e</sub> class have been investigated by several. An innovative design including heat pipes between a SOFC stack and an allothermal gasifier is described in [1]. Fryda et al. [2] studies the performance of a CHP system of less than 1MW<sub>e</sub> and consisting of an autothermal gasifier combined with a MGT and/or SOFC.

This study focus on the performance of a system combining an up scaled version (~500kW<sub>th</sub>) of the two-stage gasifier named Viking and a SOFC or a SOFC-MGT system. Viking is a 75kW<sub>th</sub> autothermal (air blown) fixed bed biomass gasifier demonstrated at the Technical University of Denmark and it is described in detail in [3]. The Viking gasifier produces almost no tars, which is favourable for downstream SOFC operation. Hofmann et al. [4] has operated a SOFC on cleaned syngas from the Viking gasifier for 150 hours without degradation.

The present study is based on zero dimensional and steady-state modelling in the simulation tool DNA [5]. DNA has incorporated thermodynamic property data, is component based and is developed at The Technical University of Denmark.

## 2 System description

Two different combined heat and power systems are investigated in this study, both based on syngas production from an up scaled Viking gasifier. A flow sheet of the two systems is depicted in Figure 1. The modelled gasifier system is slightly simplified, but aims at the same resulting gas composition and cold gas efficiency as for the Viking gasifier. In the gasifier model the dryer is heated by hot syngas. The steam production from the dryer is added to the preheated air and dry wood together with mixed air and steam are fed to the gasifier. The raw product gas is cooled to 90°C in three steps; air preheating, wood drying and syngas cooling producing hot water for district heating. The cooled syngas is then cleaned from impurities as particles and sulphur compounds before some of the water in the gas is condensed through cooling to 50°C. The cleaned and partly dried syngas is then converted into electricity and heat in a bottoming cycle consisting of a SOFC or both a SOFC and a MGT. These two system configurations will from now on be referred as the Gasifier-SOFC and the Gasifier-SOFC-MGT configuration, respectively. In the Gasifier-SOFC-MGT configuration all the components in the flow sheet are in use. With respect to Figure 1 the recuperator and gas turbine are bypassed in the Gasifier-SOFC arrangement, thus the syngas and air compressors work as blowers due to no pressurization. In addition the syngas compressor works as a roots blower for the gasifier system and not illustrated is a generator. In the Gasifier-SOFC configuration the syngas and air blowers are driven by an electric motor.



The gasifier component calculates the produced syngas composition as well as the produced ashes based on the inlet media composition and the operating conditions. The input parameters defining the operating conditions for the gasifier submodel are given in Table 1. The gasifier pressure loss is defined as the difference between the inlet air and steam mixture and the outlet syngas.

*Table 1: Inputs to the gasifier submodel*

### 3.1 Gasifier model validation

Risø-R-1712(EN)

Wood chips from beech with small amounts of oak are used in the model as for the Viking gasifier reported in Ahrenfeldt et al. [3].

As seen in Table 2 the produced syngas composition and the lower heating value (LHV) from the gasifier model is close to the Viking data. The overall performance of the modelled gasifier is also similar to the Viking gasifier as expressed in the cold gas efficiencies.

	Viking [3]	Gasifier model
H <sub>2</sub> (vol-%)	30.5	29.9
CO (vol-%)	19.6	20.8
CO <sub>2</sub> (vol-%)	15.4	13.5
CH <sub>4</sub> (vol-%)	1.16	1.19
N <sub>2</sub> (vol-%)	33.3	34.2
LHV (MJ/kg)	6.2	6.3
Cold gas eff.	93%	94%

Table 2: Dry syngas composition, lower heating value as well as cold gas efficiency for the Viking gasifier and the modelled gasifier, respectively

## 4 Solid Oxide Fuel Cell model

The SOFC stack component calculates the air and fuel outlet compositions as well as the power production. The calculations are based on the inlet air and fuel compositions and flow rates as well as the other operating conditions of the SOFC. The SOFC submodel includes an electrochemical model for predicting the performance of the SOFC. The operating conditions are partly described by input parameters given to the SOFC submodel and these are presented in Table 3.

Fuel utilization factor	$UF$	0.85
Operating temperature	$T_{SOFC}$	800°C
Anode pressure loss	$\Delta p_a$	5 mbar
Cathode pressure loss	$\Delta p_c$	10 mbar
Current density	$i$	300 mA cm <sup>-2</sup>

Table 3: Inputs to the SOFC submodel

In the submodel only H<sub>2</sub> is electrochemically converted in the SOFC anode, but the model takes into account that CO produces an extra H<sub>2</sub> through the water-gas-shift (WGS) reaction, while four additional H<sub>2</sub> molecules are produced from CH<sub>4</sub> through internal steam reforming and WGS of produced CO (full conversion is assumed). The total mole flow of H<sub>2</sub> on the anode after internal steam reforming and WGS is expressed in Eq. (1).

$$\dot{n}_{H_2, \text{tot}} = \dot{n}_{H_2, \text{in}} + \dot{n}_{CO, \text{in}} + 4\dot{n}_{CH_4, \text{in}} \quad (1)$$



The amount of hydrogen that is converted depends on the fuel utilization factor ( $UF$ ) and this amount is electrochemically converted in the anode. The electrode reactions and the overall fuel cell reaction are as shown in Eq. (2) to (4).

From the overall fuel cell reaction it is seen that the amount of consumed oxygen is half the amount of consumed hydrogen. The cathode outlet composition can then be found by

the following equations if the only species taking into account are O<sub>2</sub>, N<sub>2</sub>, CO<sub>2</sub>, H<sub>2</sub>O and Ar.

$$\dot{n}_{O_2, \text{con}} = \frac{UF \dot{n}_{H_2, \text{in}}}{2} \quad (5)$$

$$\dot{n}_{c, \text{out}} = \dot{n}_{c, \text{in}} - \dot{n}_{O_2, \text{con}} \quad (6)$$

$$y_{O_2, \text{out}} = \frac{\dot{n}_{c, \text{in}} y_{O_2, \text{in}} - \dot{n}_{O_2, \text{con}}}{\dot{n}_{c, \text{out}}} \quad (7)$$

$$y_{j, \text{out}} = \frac{\dot{n}_{c, \text{in}} y_{j, \text{in}}}{\dot{n}_{c, \text{out}}}, j = \{N_2, CO_2, H_2O\} \quad (8)$$

$$y_{Ar, \text{out}} = 1 - y_{O_2, \text{out}} - y_{N_2, \text{out}} - y_{CO_2, \text{out}} - y_{H_2O, \text{out}} \quad (9)$$

The fuel composition leaving the anode is calculated by the Gibbs minimization method [6] as described for the gasifier submodel. Equilibrium at the anode outlet temperature and pressure is assumed for the following species: H<sub>2</sub>, CO, CO<sub>2</sub>, H<sub>2</sub>O, CH<sub>4</sub> and N<sub>2</sub>. The equilibrium assumption is fair since the methane content in this study is low enough for such kind of assumption to be made. The heat consumed by the endothermic internal reforming reactions is taken into account by the Gibbs minimization method. More internal reforming means more cooling of the SOFC.

The power production from the SOFC depends on the amount of chemical energy fed to the anode, the reversible efficiency ( $\eta_{\text{rev}}$ ), the voltage efficiency ( $\eta_v$ ) and the fuel utilization factor ( $UF$ ). It is defined in mathematical form in Eq. (10).

$$P_{\text{SOFC}} = [LHV_{H_2} \dot{n}_{H_2, \text{in}} + LHV_{CO} \dot{n}_{CO, \text{in}} + LHV_{CH_4} \dot{n}_{CH_4, \text{in}}] \eta_{\text{rev}} \eta_v UF \quad (10)$$

The reversible efficiency is the maximum possible efficiency defined as the relationship between the maximum electrical energy available (change in Gibbs free energy) and the fuels LHV. This is shown in Eq. (11) and the definition of the change in Gibbs free energy is shown in Eq. (12). The voltage efficiency express the electrochemical performance of the SOFC and the calculation of the voltage efficiency is described in the following subsection.

$$\eta_{\text{rev}} = \frac{(A\bar{g}_f)_{\text{fuel}}}{LHV_{\text{fuel}}} \quad (11)$$

$$\begin{aligned} (A\bar{g}_f)_{\text{fuel}} = & [(\bar{g}_f)_{H_2O} - (\bar{g}_f)_{H_2} - \frac{1}{2}(\bar{g}_f)_{O_2}] y_{H_2, \text{in}} \\ & + [(\bar{g}_f)_{CO_2} - (\bar{g}_f)_{CO} - \frac{1}{2}(\bar{g}_f)_{O_2}] y_{CO, \text{in}} \\ & + [(\bar{g}_f)_{CO_2} + 2(\bar{g}_f)_{H_2O} - (\bar{g}_f)_{CH_4} - 2(\bar{g}_f)_{O_2}] y_{CH_4, \text{in}} \end{aligned} \quad (12)$$

#### 4.1 Electrochemical model

The electrochemical model is used to calculate the cell potential and the voltage efficiency of the SOFC. Both depend on the operating conditions such as temperature, pressure, gas compositions, fuel utilization and load (current density). The cell potential and voltage efficiency is defined in Eq. (13) and (14), respectively.

$$V_{\text{cell}} = E - V_{\text{act}} - V_{\text{ohm}} \quad (13)$$

$$\eta_v = \frac{V_{\text{cell}}}{E} \quad (14)$$

In the following the reversible open circuit voltage ( $E$ ), the activation overpotential ( $V_{\text{act}}$ ) and the ohmic overpotential ( $V_{\text{ohm}}$ ) are calculated. Traditionally a concentration overpotential term is included in Eq. (13). The concentration overpotential is a result of the limitations of transporting the reactants to the active cell area. In Larminie et al. [7] it is described as a voltage drop caused by the pressure change associated with the consumption of reactants. As a result of the current being drawn from the cell, the average partial pressure of reactants is lower than at the inlet. Thus, in this study the concentration overvoltage is taken into account by using average partial pressures when calculating  $E$  and  $V_{\text{act}}$ .

$E$  can be calculated from the Nernst equation:

$$E = \frac{-\Delta \bar{g}_f^0}{n_e F} + \frac{RT}{n_e F} \ln \left( \frac{\bar{p}_{\text{H}_2, \text{tot}} \sqrt{\bar{p}_{\text{O}_2}}}{\bar{p}_{\text{H}_2\text{O}}} \right) \quad (15)$$

Since it is assumed that all CO and CH<sub>4</sub> are converted to H<sub>2</sub> before the electrochemical reactions take place, the change in standard Gibbs free energy ( $\Delta \bar{g}_f^0$ ) is and the number of electrons transferred for each molecule of fuel ( $n_e$ ) are determined for the reaction of H<sub>2</sub> only. Thus,  $n_e = 2$  and  $\Delta \bar{g}_f^0 = (\bar{g}_f^0)_{\text{H}_2\text{O}} - (\bar{g}_f^0)_{\text{H}_2} - \frac{1}{2} (\bar{g}_f^0)_{\text{O}_2}$ . The partial pressure of species  $j$  is an average across the respective electrode and is here defined as an arithmetic mean between inlet and outlet as shown in Eq. (16) and (17). The average partial pressure of available hydrogen after internal steam reforming and WGS of CH<sub>4</sub> and CO can be determined from the overall steam reforming and WGS reaction including all species. It is defined in Eq. (18).

$$\bar{p}_j = \left( \frac{y_{j, \text{out}} - y_{j, \text{in}}}{2} \right) p_a, \quad j = \{\text{H}_2, \text{CO}, \text{CH}_4, \text{CO}_2, \text{H}_2\text{O}, \text{N}_2\} \quad (16)$$

$$\bar{p}_{\text{O}_2} = \left( \frac{y_{\text{O}_2, \text{out}} - y_{\text{O}_2, \text{in}}}{2} \right) p_c \quad (17)$$

$$\bar{p}_{\text{H}_2, \text{tot}} = \left( \frac{\bar{p}_{\text{H}_2} + \bar{p}_{\text{CO}} + 4\bar{p}_{\text{CH}_4}}{\bar{p}_{\text{H}_2} + \bar{p}_{\text{CO}} + 3\bar{p}_{\text{CH}_4} + \bar{p}_{\text{CO}_2} + \bar{p}_{\text{H}_2\text{O}} + \bar{p}_{\text{N}_2}} \right) p_a \quad (18)$$

The activation overpotential is due to an energy barrier (activation energy) that the reactants must overcome in order to drive the electrochemical reactions. The activation overpotential is non-linear and is dominant at low current densities ( $i$ ). The activation overpotential is defined as (cf. [8]):

$$V_{\text{act}} = V_{\text{act}, a} + V_{\text{act}, c} = \frac{2RT}{n_e F} \left[ \sinh^{-1} \left( \frac{i + i_n}{2i_{0, a}} \right) + \sinh^{-1} \left( \frac{i + i_n}{2i_{0, c}} \right) \right] \quad (19)$$

The internal current density ( $i_n$ ) is added to the actual fuel cell current density in order to account for the mixed potential caused by fuel crossover. The importance of the internal current density in the case of SOFCs is much less than for low temperature fuel cells and the value of  $i_n$  is usually very small [7]. The exchange current density ( $i_0$ ) is a measure of the level of activity on the electrode at  $i=0$  mA cm<sup>-2</sup> and is defined as (cf. [9]):

$$i_{0, a} = 2.13 \times 10^7 \left( \frac{\bar{p}_{\text{H}_2, \text{tot}} \bar{p}_{\text{H}_2\text{O}}}{p_a^2} \right) \exp \left( \frac{-110000}{RT} \right) \quad (20)$$

$$i_{0, c} = 1.49 \times 10^7 \left( \frac{\bar{p}_{\text{O}_2}}{p_c} \right)^{0.25} \exp \left( \frac{-110000}{RT} \right) \quad (21)$$

The ohmic overpotential is caused by the electrical resistance for the ions passing through the electrolyte as well as for the electrons passing through the electrodes and interconnects. The ohmic overpotential is defined below (cf. [9] and [12]).

$$V_{\text{ohm}} = i ASR \quad (22)$$

$$ASR = ASR_a + ASR_c + ASR_e + ASR_i \quad (23)$$

$$ASR_j = \delta_j a_{\text{ohm},j} \exp\left(\frac{b_{\text{ohm},j}}{T}\right), j = \{a, c, e, i\} \quad (24)$$

The thicknesses of the different layers ( $\delta$ ) and the constants  $a_{\text{ohm}}$  and  $b_{\text{ohm}}$  used are listed in Table 4.

$R$	$8.314 \text{ J K}^{-1} \text{ mol}^{-1}$	
$F$	$96485 \text{ C mol}^{-1}$	
$n_e$	2	
$i_n$	$2 \text{ mA cm}^{-2}$	[9]
$\delta_a$	$750 \times 10^{-4} \text{ cm}$	[10]
$\delta_c$	$50 \times 10^{-4} \text{ cm}$	[10]
$\delta_e$	$40 \times 10^{-4} \text{ cm}$	[10]
$\delta_i$	$100 \times 10^{-4} \text{ cm}$	[11]
$a_{\text{ohm},a}$	$0.00298 \times 10^{-3} \text{ k}\Omega\text{cm}$	[12]
$b_{\text{ohm},a}$	-1392 K	[12]
$a_{\text{ohm},c}$	$0.00811 \times 10^{-3} \text{ k}\Omega\text{cm}$	[12]
$b_{\text{ohm},c}$	600 K	[12]
$a_{\text{ohm},e}$	$0.00294 \times 10^{-3} \text{ k}\Omega\text{cm}$	[12]
$b_{\text{ohm},e}$	10350 K	[12]
$a_{\text{ohm},i}$	$0.1256 \times 10^{-3} \text{ k}\Omega\text{cm}$	[12]
$b_{\text{ohm},i}$	4690 K	[12]

Table 4: Inputs for the electrochemical model

## 4.2 Electrochemical model calibration

The described electrochemical model has been calibrated against experimental data, see Figure 2.

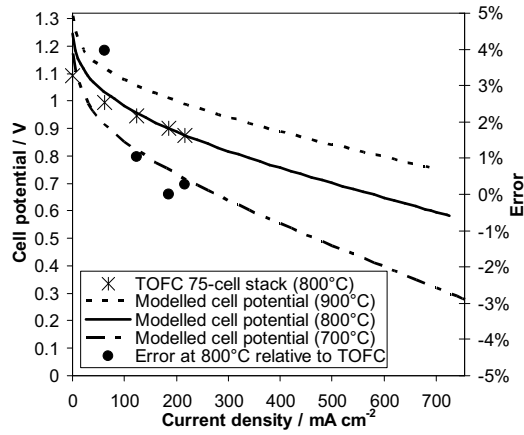


Figure 2: Single cell polarization curves based on a 75-cell stack and the SOFC model, respectively

Since the model aims at the performance of 2<sup>nd</sup> generation SOFCs from Topsoe Fuel Cell A/S (TOFC) and Risø National Laboratory, published data for this SOFC type has been used. The  $ASR$  has been calibrated against a value of  $0.15 \Omega \text{ cm}^2$  at  $850^\circ\text{C}$  as published by [13] and the resulting cell potential has been calibrated against a polarization curve



(75-cell stack,  $12 \times 12 \text{ cm}^2$ ,  $800^\circ\text{C}$  and fuelled with  $\text{H}_2$  and  $\text{N}_2$ ) published by [14]. An active cell area of  $81 \text{ cm}^2$  has been assumed. Both modelled and experimental data as well as the error relative to the experimental data are presented in Figure 2.

The model shows excellent agreement with the experimental data above a current density of  $100 \text{ mA cm}^{-2}$ . The current density of  $300 \text{ mA cm}^{-2}$  is chosen to represent the SOFC load in the following results.

## 5 Peripheral equipment

Modelling of peripheral components like compressors, turbines and heat exchangers are standard and therefore not described in detail.

The throughput of wet biomass is  $154.8 \text{ kg h}^{-1}$  (corresponds to  $499.2 \text{ kW}_{\text{th}}$  (LHV)). Thus it is assumed that the Viking gasifier can be scaled up from a nominal  $\sim 75 \text{ kW}_{\text{th}}$  [3]. The biomass dryer reduces the water content in the biomass from 32.2 wt-% to 5 wt-% by heating it to  $150^\circ\text{C}$  and the air for the gasifier is preheated to  $780^\circ\text{C}$  by the hot product gas.

The inlet temperature to the SOFC anode and cathode are maintained at  $150^\circ\text{C}$  and  $200^\circ\text{C}$  below the outlet temperature, respectively.

The pressure loss in every component in the SOFC air supply stream and burner exhaust stream is assumed to be 10 mbar, while the pressure loss in each of the rest of the components is assumed to be 5 mbar, except the burner that has a pressure loss of 0.6% (equals 1.5 mbar when 2.5 bar at inlet).

The gas cleaner is a baghouse filter removing particulates and it is assumed that the cleaned syngas can be used directly in a SOFC. The condenser removes some of the water content in the syngas resulting in a content of water in the cleaned and dried syngas of 12.7 vol-%. The resulting steam to carbon ratio (S/C) is 0.41, which is somewhat low, but is justified by the very low tar content in the Viking syngas.

The isentropic and mechanical efficiency of the compressors are 75% and 98%, respectively, and the isentropic efficiency of the MGT expander is 84%. The performance of the compressors and the MGT expander are taken from Fryda et al. [2] and corresponds to common performance data for a MGT of this scale. The recuperator effectiveness is assumed to be 85% and the generator efficiency is assumed to be 99%. In the Gasifier-SOFC configuration the SOFC operating pressure is  $\sim 1$  bar and in the Gasifier-SOFC-MGT case the SOFC operating pressure is 2.5 bar (this pressure is varied in the results section).

No heat losses are taken into account. Introducing heat losses from the gas cleaner will only affect the heat production from the condenser since the temperature after the condenser is fixed to  $50^\circ\text{C}$ .

The outlet pressure from the MGT depends on the total pressure loss downstream the MGT, since it is the exhaust pressure which is fixed to 1.013 bar. Because of the recuperator and exhaust cooler the outlet pressure from the MGT is 1.033 bar. The district heating (DH) water is assumed to be  $30^\circ\text{C}$  at inlet and  $80^\circ\text{C}$  at outlet.

## 6 Results and discussion

In the following results the inputs presented in the previous sections are used unless something else is stated. The system configurations are previously described in detail.

The performance of the different system configurations vary greatly with the operating conditions and namely the operating pressure of the SOFC (in the Gasifier-SOFC-MGT case) are of great importance to the resulting system performance. The Gasifier-SOFC-MGT configuration has an optimum with regard to its operating pressure, while the

Gasifier-SOFC arrangement always operates at atmospheric pressure – illustrated in Figure 3. The Gasifier-SOFC configuration performs an electric efficiency of 36.4%. By combining the SOFC and MGT in the Gasifier-SOFC-MGT configuration the electrical efficiency reaches 50.3% at an optimum operating pressure of 2.5 bar. This is a substantial increase in efficiency caused by the utilization of unconverted fuel from the SOFC (fuel utilization of 85%) in the MGT as well as the pressurized operation of the SOFC. In the Gasifier-SOFC-MGT case the turbine inlet temperature (TIT) is varying with the SOFC operating pressure and has a value of 697°C at 2.5 bar. It is the recuperator that ensures an optimum at a relatively low operating pressure.

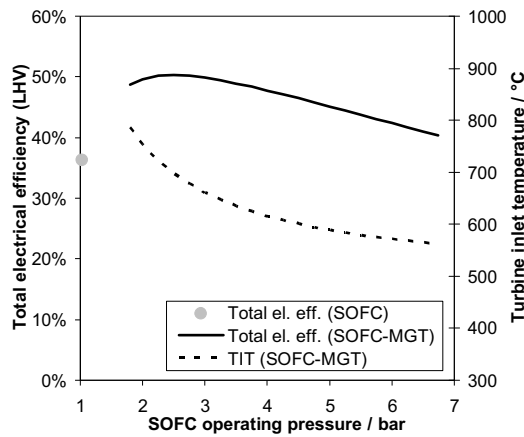


Figure 3: Electric efficiency and TIT at different SOFC operating pressures

The performance of both system arrangements strongly depend on the SOFC operating temperature as depicted in Figure 4. Decreasing the temperature by 100°C to 700°C lowers the electrical efficiency to 28.8% and 44.4% in the Gasifier-SOFC and Gasifier-SOFC-MGT case, respectively. This corresponds to a drop of 7.6 and 5.9 percentage points, respectively. The research and development working on lowering the SOFC operating temperature in order to use cheaper materials will influence the system performance presented here and potentially other bottoming cycles could be beneficial, e.g. a Rankine cycle.

The sensitivity of the model results to the chosen SOFC current density is shown in Figure 5.

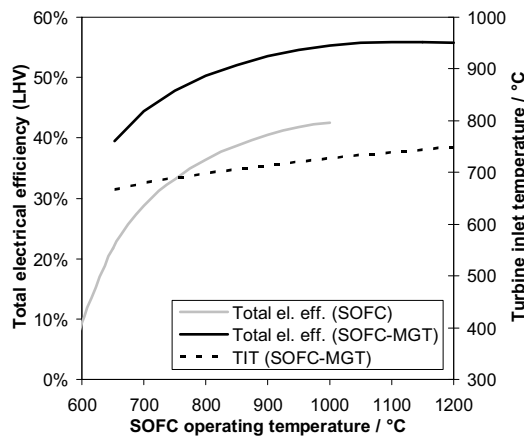


Figure 4: Electric efficiency and TIT at different SOFC operating temperatures

At the reference current density value of  $300 \text{ mA cm}^{-2}$  the SOFC voltage efficiency is 39.6% in the Gasifier-SOFC arrangement and 40.8% in the Gasifier-SOFC-MGT case. The difference is due to the pressure. Raising the SOFC load to  $500 \text{ mA cm}^{-2}$  reduces the

voltage efficiency (defined in Eq. (14)) to 34.6% and 35.7% in the Gasifier-SOFC and Gasifier-SOFC-MGT cases, respectively, meaning a reduction in the total electrical efficiency to 31.5% and 46.7% - a drop of 4.9 and 3.6 percentage points. This is a relative change in electrical efficiency of 13.5% and 7.2%, respectively, for a 66.7% increase in current density.

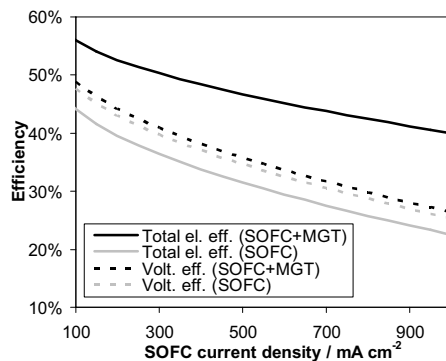


Figure 5: Electrical efficiency and SOFC voltage efficiency as a function of SOFC current density

Key data for the two system configurations studied are presented in Table 5 based on the reference input values presented in the previous sections. The Gasifier-SOFC-MGT configuration clearly has the best electrical efficiency, while the CHP efficiencies do not differ significantly. In the Gasifier-SOFC-MGT case, the power production is mainly from the SOFC producing 76.4% of the power. The exact values of the efficiencies will be slightly lower when incorporating heat losses, a more accurate efficiency of the gasifier system and possible more extensive gas cleaning, but the comparison of the systems performance is still valid.

		Gasifier -SOFC	Gasifier -SOFC-MGT
Biomass input	/ kg h <sup>-1</sup>	154.8	154.8
	/ kW <sub>th,LHV</sub>	499.2	499.2
$p_{\text{SOFC}}$ / bar		1.034	2.5
$P_{\text{MGT,net}}$ / kW <sub>el</sub>		-	59.2
$P_{\text{SOFC,net}}$ / kW <sub>el</sub>		181.5	191.8
$P_{\text{total,net}}$ / kW <sub>el</sub>		181.5	251.0
DH production / kJ s <sup>-1</sup>		216.6	146.7
$\eta_{\text{el}}$ / % (LHV)		36.4	50.3
$\eta_{\text{CHP}}$ / % (LHV)		79.74	79.68

Table 5: Key data for the studied systems

## 7 Conclusion

A study on the system performance of an up scaled Viking gasifier (~500 kW<sub>th</sub>) with either a downstream SOFC or SOFC-MGT arrangement has been conducted by zero dimensional process modelling. A SOFC submodel has been developed including an electrochemical model predicting the SOFC performance at different operating conditions. This submodel has been calibrated against published TOFC stack performance data. The reference conditions for the SOFC has been an operating temperature of 800°C, a fuel utilization of 85% and a current density of 300 mA cm<sup>-2</sup>. The optimal operating SOFC-MGT pressure has been found to be 2.5 bar, while the SOFC without MGT operated at atmospheric pressure. The MGT utilized the unconverted syngas from the SOFC to produce more power as well as pressurizing the SOFC bettering the electrical efficiency compared to operation with the SOFC alone -

from  $\eta_{el}=36.4\%$  to  $\eta_{el}=50.3\%$ . These efficiencies were very sensitive to the SOFC operating temperature, while only a moderate sensitivity to the SOFC current density was observed.

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